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Attorney Docket No.: 10004.204-US

PATENT

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

In re Application of: Andersen et al.

Confirmation No: 2881

Serial No.: 09/925,576

Group Art Unit: To be assigned

Filed: August 9, 2001

Examiner: To be assigned

For: Amylase Variants

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(name of person mailing paper)

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**RESPONSE TO NOTICE TO COMPLY WITH SEQUENCE REQUIREMENTS**

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Sir:

This paper is in response to the Notice to Comply with Sequence Requirements dated March 25, 2002. The Notice stated that the computer readable form that was filed on December 13, 2001 was unreadable.

Applicants therefore enclose another computer readable form of the Sequence Listing. The content of the paper copy that was filed with the application and of the enclosed computer readable form is the same. This submission contains no new matter.

The Examiner is hereby invited to contact the undersigned by telephone if there are any questions concerning this amendment or application.

Respectfully submitted,

Date: May 24, 2002

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APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
09/925,576	08/09/2001	Carsten Andersen	10004.204-US

CONFIRMATION NO. 2881

25908  
NOVOZYMES NORTH AMERICA, INC.  
C/O NOVO NORDISK OF NORTH AMERICA, INC.  
405 LEXINGTON AVENUE, SUITE 6400  
NEW YORK, NY 10174

## FORMALITIES LETTER



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Date Mailed: 03/25/2002

### NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant is given **TWO MONTHS FROM THE DATE OF THIS NOTICE** within which to file the items indicated below to avoid abandonment. Extensions of time may be obtained under the provisions of 37 CFR 1.136(a).

- A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing." Applicant must provide a substitute computer readable form (CRF) copy of the "Sequence Listing" and a statement that the content of the sequence listing information recorded in computer readable form is identical to the written (on paper or compact disc) sequence listing and, where applicable, includes no new matter, as required by 37 CFR 1.821(e), 1.821(f), 1.821(g), 1.825(b), or 1.825(d).

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